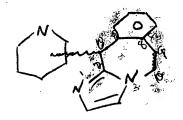
ONLINE SEARCH REQUEST FORM

********	**********
USER DATION	SERIAL NUMBER US/142474
ART UNIT 12M	PHONE 74710 DATE

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You may include a copy of the broadest and or relevant claim(s).



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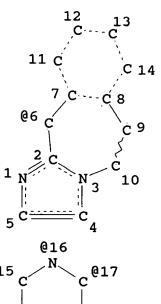
___CAS ONLINE

____DARC/QUESTEL

___ DIALOG ___ SDC

____ OTHER

=> d 17 que stat L3 STR



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VPA 6-16/17/18/19 U NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 20

STEREO ATTRIBUTES: NONE

L5 234 SEA FILE=REGISTRY SSS FUL L3 L6 STR

NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED **GRAPH ATTRIBUTES:**

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RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 20

STEREO ATTRIBUTES: NONE

L7 234 SEA FILE=REGISTRY SUB=L5 SSS FUL L6

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234 ANSWERS

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L8 2 L7

=> d 1-2 an .mh

L8 ANSWER 1 OF 2 CA COPYRIGHT 1994 ACS

AN 119:8809 CA

'TI Preparation of imidazo[2,1-b][3]benzazepine derivatives as antiallergy agents

SO Eur. Pat. Appl., 66 pp.

CODEN: EPXXDW

IN Janssens, Frans Eduard; Diels, GAston Stanislas Marcella; Leenaerts,
Joseph Elisabeth

PI EP 518435 A1 921216

AI EP 92-201666 920609

PY 1992

GI

- Title compds. I (R1 = H, halo, C1-4 alkyl, C1-4 alkoxy; R2 = R1; R3 AΒ = H, (substituted) C1-4 alkyl, substituted ethenyl; R4 = H, (HO)-C1-4-alkyl, Ph, halo; R5 = H, C1-4 alkyl, halo; L = H, (substituted) C1-6 alkyl;, C1-4 alkoxycarbonyl, HO2C-C1-4-alkoxy, aryl, aryloxy, arylcarbonyl, (substituted) C3-6 alkenyl, etc.), isomer or salt thereof, useful as antiallergy agents (no data), are 3-FC6H4CH2CH2OH, MeCH2NEt2, and CH2Cl2 were added to MeSO2Cl to give 3-FC6H4CH2CH2OSO2Me which was treated with 1H-imidazole, H2CO3 and THF to give 1-(2-fluorophenyl)ethyl-1H-imidazole (II). MeCH2CH2NHCHMe2 and THF under N was added BuLi, II and Et 1-methyl-4-piperidinecarboxylate to give the methanone deriv. which was heated with AlCl3 and NaCl at 120-140.degree. to give title compd. III. A large no. of I were prepd.
- L8ANSWER 2 OF 2 CA COPYRIGHT 1994 ACS
- AN 117:90284 CA
- TI Preparation of substituted imidazobenzazepines and imidazopyridoazepines as platelet activating factor antagonists
- SO PCT Int. Appl., 47 pp. CODEN: PIXXD2
- IN Friary, Richard J.
- PI WO 9206981 Al 920430
- ΑI WO 91-US7156 911004
- PY 1992

GI

$$R^{2}$$
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AB Title compds. [I; R1 = H, (cyclo)alkyl, CF3, (cyclo)alkoxy, (substituted) (hetero)aryl; R2, R3 = H, (cycloalkyl), CF3, NO2, halo, OR7, NR8R9, S OmR10; m = 0-2; R4 = H, alkyl, (substituted) arylmethyl; R5, R6 = H, (cyclo)alkyl, (substituted) (hetero)aryl, arylmethyl; R5R6 = (CH2)k; k = 3-5; R7-R9 = H, alkyl, alkylcarbonyl, (hetero)arylcarbonyl; R10 = (cyclo)alkyl, (substituted) (hetero)aryl, arylmethyl; n = 0-3; Z = 0,S; Q = CH, N, NO; Z .noteq. S when Q = NO; one of the dotted lines = double bond], were prepd. Thus, 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride, 6,11-dihydro-11-(4-piperidinylidene)-5H-imidazo[2,1-b][3]benzazepine (prepn. given), pyridine-4-carboxylic acid N-oxide, and 1-hydroxybenzotriazole were stirred at 0 to 25.degree. in CH2Cl2 overnight to give title compd. II. II inhibited PAF-induced platelet aggregation with IC50 = 5 .mu.M. Tablets were prepd. contg. II.

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=> s 18 L10 0 L7

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L1 STR
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L3 STR L1
L4 14 S L3
L5 234 S L3 FUL
L6 STR L3

L7 234 SEARCH L6 SUB=L5 FUL

FILE 'CA' ENTERED AT 10:25:49 ON 10 AUG 94
L8 2 S L7

FILE 'CAOLD' ENTERED AT 10:26:29 ON 10 AUG 94

L9 0 S LL8 L10 0 S L8

FILE 'CAPREVIEWS' ENTERED AT 10:26:44 ON 10 AUG 94 L11 0 S L7

FILE 'REGISTRY' ENTERED AT 10:26:53 ON 10 AUG 94

=> save 17 datlow142/a
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